

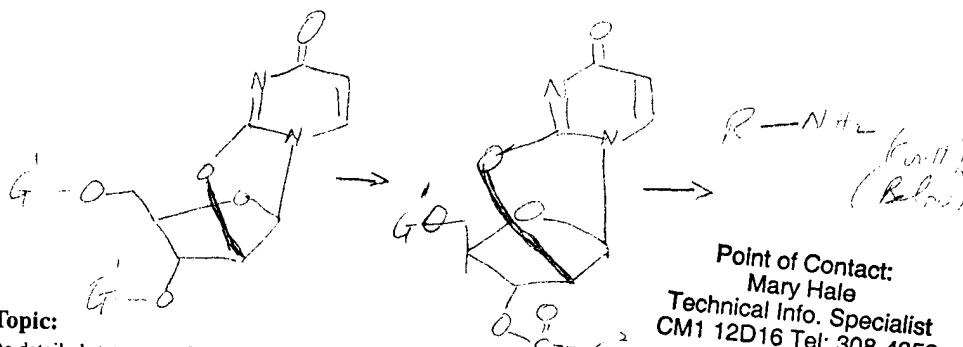
SEARCH REQUEST FORM

7 290

Examiner # (Mandatory): L. E. Crane Requester's Full Name: sameArt Unit 1623 Location (Bldg/Room#): 8D-14 Phone (circle 305 306 308) 4639Serial Number: 09092,167 Results Format Preferred (circle): PAPER DISK E-MAILTitle of Invention See copy of claims attached And particularly copies of
Figures attachedInventors (please provide full names): " "Earliest Priority Date: 11/13/98

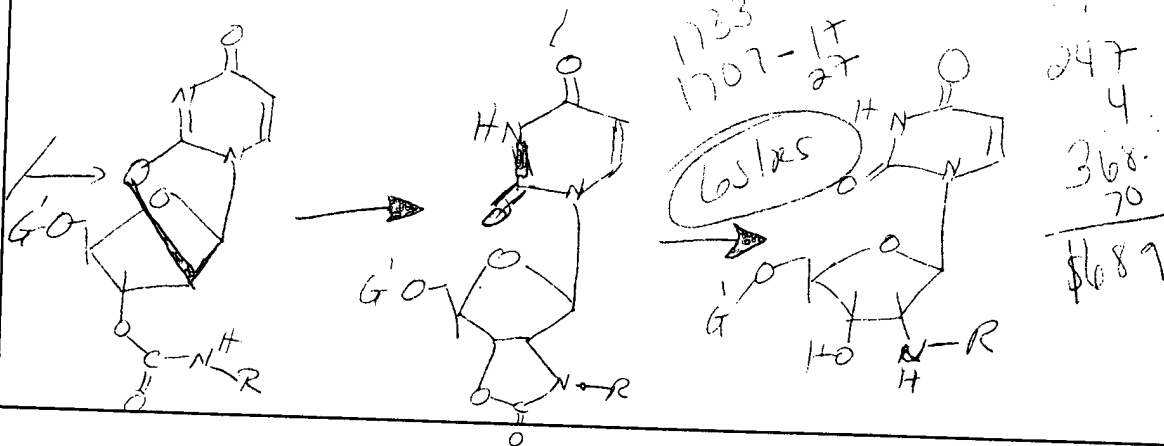
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CLAIMS

We claim:

1. A method for making a modified nucleoside comprising a covalently attached signalling moiety or signalling moiety precursor, said method comprising:

- a) adding an anhydro-nucleoside and a signalling moiety or signalling moiety precursor comprising a primary amine in the presence of an activation agent to form an activated anhydro-nucleoside;
- b) treating said activated anhydro-nucleoside with a cyclization agent to form a cyclized intermediate; and
- c) treating said cyclized intermediate with a base to form said modified nucleoside.

2. A method according to claim 1 further comprising adding a phosphoramidite group to said modified nucleoside.

3. A method according to claim 2 further comprising incorporating said phosphoramidite modified nucleoside into a growing nucleic acid.

4. A method according to claim 1 wherein said nucleoside is a naturally occurring nucleoside.

5. A method according to claim 1 wherein said nucleoside is a nucleoside analog.

6. A method according to claim 1 wherein said activating agent is carbonyldimidazole.

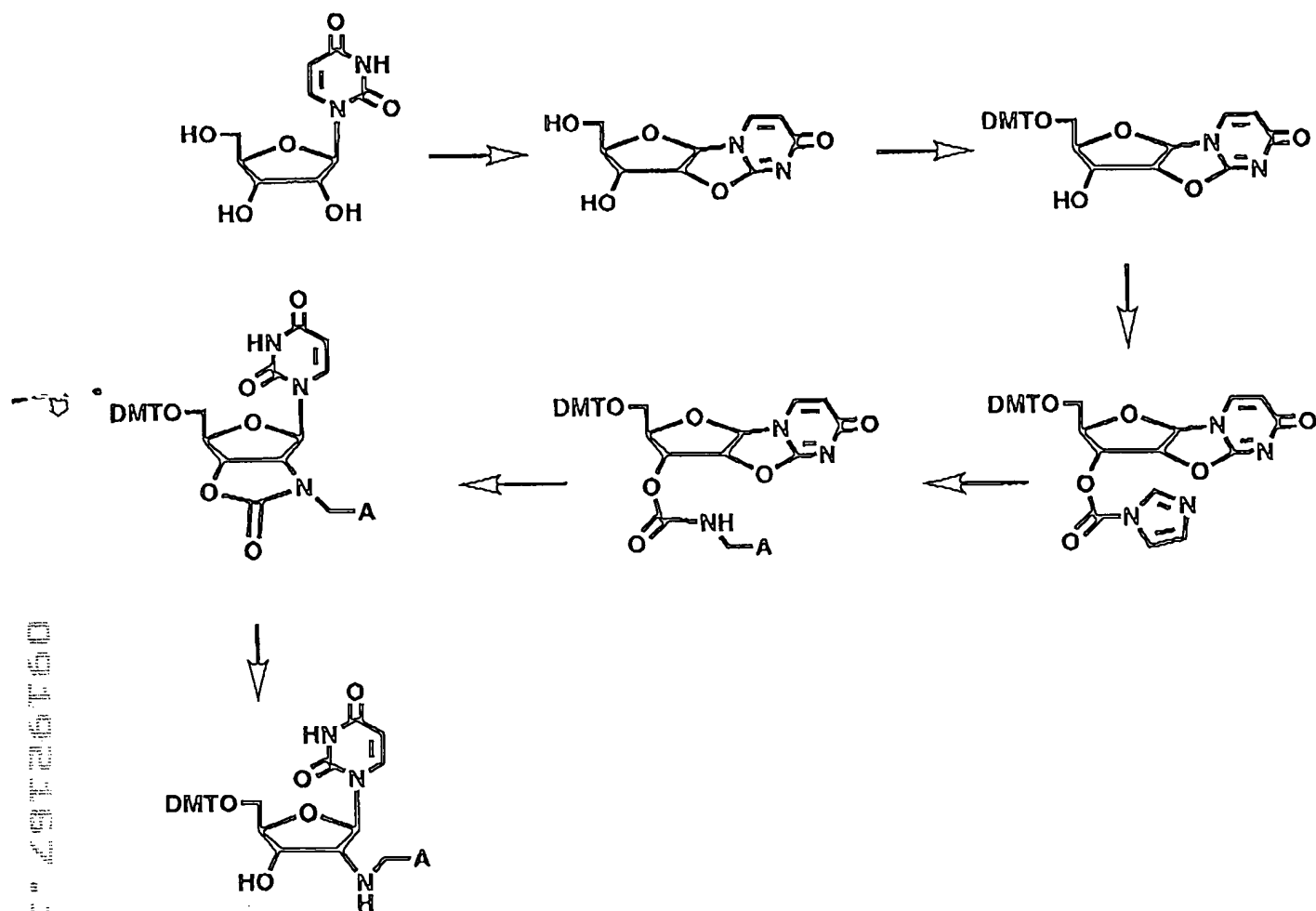
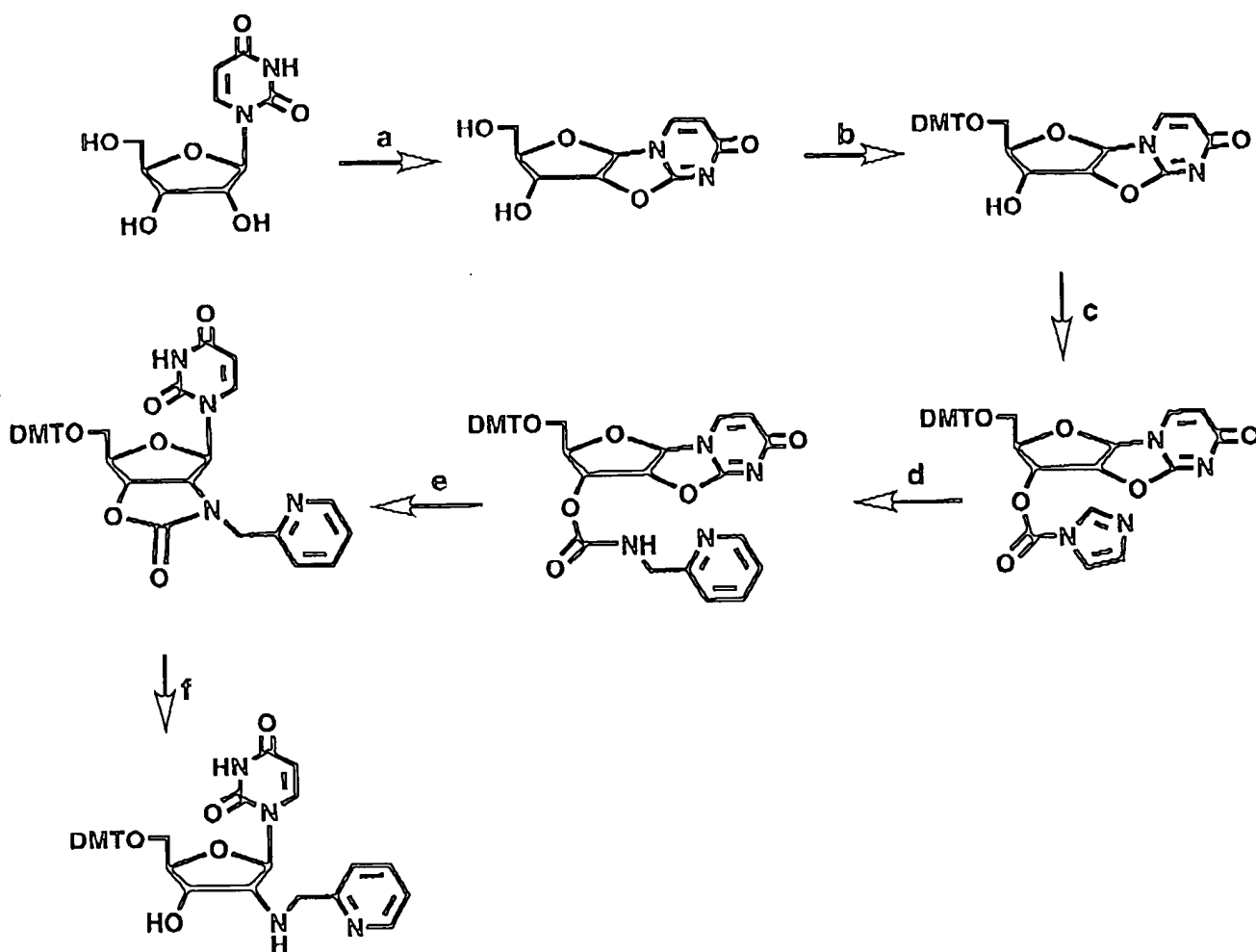


Fig 1



a) Diphenylcarbonate, DMF, 110° C, 8 hours; b) DMTCl, cat. DMAP, CH_2Cl_2 , 25°C, 24 hrs.; c) 1,1'-Carbonylimidazole, CH_2Cl_2 , 24 hrs.; d) 2-aminomethylpyridine, DIEA, CH_2Cl_2 , 24 hrs.; e) DBU, THF, 48 hrs. 25°C; f) NaOH / MeOH / H_2O , 24 hr, 25°C.

Fig ②

J.D.

Synthesis of A Metallated Phosphoramidate

Fig 3

